Brain volume, asymmetry and intellectual impairment in relation to sex in early-onset schizophrenia

SIMON L. COLLINSON, CLARE E. MACKAY, ANTHONY C. JAMES, DIGBY J. QUESTED, TANIA PHILLIPS, NEIL ROBERTS and TIMOTHY J. CROW

Background  Accumulating evidence suggests that early-onset schizophrenia arises from a disturbance in the normal trajectory of cerebral development.

Aims  To investigate brain structure, asymmetry and IQ in early-onset schizophrenia.

Method  Volumes of left and right cerebral hemispheres and IQ were assessed in 33 participants with early-onset DSM–IV schizophrenia and 30 members of a matched, normal control group.

Results  Total brain volume was significantly smaller in the group with early-onset disease (‘cases’) relative to the control group (4.5%), especially for the left hemisphere in males (6.0%). A significant sex × diagnosis interaction in hemisphere asymmetry revealed that the female cases group had significantly reduced rightward asymmetry relative to the female control group and that the male cases tended to have reduced leftward asymmetry relative to the male control group. Decreased left hemisphere volume in males and decreased rightward hemispheric asymmetry in females correlated with reduced IQ.

Conclusions  Sexually dimorphic alterations in asymmetry correlate with degree of intellectual impairment in early-onset schizophrenia.

Declaration of interest  None. Funding details in Acknowledgements.

It has been suggested that earlier onset of psychosis reflects a greater deviation than later-onset psychosis from the normal sexually dimorphic course of brain development leading to cerebral lateralisation (Crow et al, 1989; Crow, 1990). Cerebral hemisphere volume, as a marker for structural change, has been reported to be reduced by 5–10% in patients with early disease onset relative to age-matched controls (Table 1), compared with average reductions of 2–3% in adult schizophrenia (Harrison, 1999; Wright et al, 2000). Correspondingly, full-scale IQ scores, as a marker for functional change, have been reported to be 70–95 in early-onset samples (Table 2) relative to 85–100 in adult patients (Aylward et al, 1984; Gold et al, 1999). Correlations between brain volume and cognitive ability in healthy individuals (Reiss et al, 1996; Wickett et al, 2000; Thompson et al, 2001) are reported to depend on age and sex (Willerman et al, 1992; Andreasen et al, 1993; Coffey et al, 1998; Raz et al, 1998; Gur et al, 1999). Here we investigate sex-dependent alterations in brain volume, asymmetry and IQ in males and females with early onset of psychosis relative to controls.

METHOD

Study group

Thirty-three (22 males, 11 females) with early-onset schizophrenia were selected from a larger sample of patients referred to the Oxford Early Psychosis Project. Patients were recruited from both in-patient and out-patient National Health Service (NHS) and private adolescent units across the south of England. All participants met DSM–IV criteria for schizophrenia (American Psychiatric Association, 1994) following a semi-structured interview using the Schedule for Affective Disorders and Schizophrenia for School-aged Children: Present and Lifetime Version (KSADS–PL; Kaufman et al, 1997); most of them had experienced only one psychotic episode. Exclusion criteria were history of significant substance misuse, brain injury, epilepsy, or other neurological or psychiatric disorder. The mean age, height, estimated age at onset, duration of illness, and medication are shown in Table 3. In addition 30 control participants (18 males, 12 females) were recruited from local general practitioners within the Oxford area. The same exclusion criteria were applied. Cases and controls were closely matched for age and had comparable levels of education. After a complete description of the study to the participants, written informed consent was obtained from both the young people and their parents.

Magnetic resonance image acquisition and analysis

Magnetic resonance images were acquired using a 1.5T Magnetom Vision whole body system (Siemens Medical System Inc., Erlangen, Germany). One hundred and fifty-six coronal T₁-weighted images were obtained using a three-dimensional spoiled gradient echo pulse sequence (time to repetition=34 ms, time to echo=9 ms, flip angle 30°). The field of view of the images was 20 cm, with 1.5 mm slice thickness. The left and right temporal lobes were optimally visualised, and their volumes best measured, on image sections oriented perpendicularly to the long axis of the hippocampus (Mackay et al, 1998). These sections were obtained by reformatting oblique sections through the acquired three-dimensional data using New Region of Interest Analysis (NRIA) software (Brain Behavior Laboratory, University of Pennsylvania, USA) running on an Ultra 10 Workstation (Sun Microsystems, California, USA), where the 256 × 256 × 156 acquired voxels of side 0.78 mm × 0.78 mm × 1.5 mm were linearly interpolated to 256 × 256 × 256 cubic voxels of side 0.78 mm. This was also a convenient sectioning direction for volume estimation of the left and right cerebral hemisphere and lateral ventricles.

Unbiased estimates of structure volume were obtained using the mathematically unbiased Cavalieri method of modern design stereology in combination with point counting (Roberts et al, 1994; Mackay et al, 1998), using EasyMeasure software (http://www.easymeasure.co.uk); see Roberts et al (2000). The posterior limit
Table 1  Demographic characteristics of study groups of patients with early-onset schizophrenia and percentage change in ventricular, temporal lobe and total cerebral volumes in early-onset cohorts relative to matched normal controls

<table>
<thead>
<tr>
<th>Study</th>
<th>n</th>
<th>Age (years)</th>
<th>Age at onset (years)</th>
<th>Total volume change (%)</th>
<th>Temporal lobe volume change (%)</th>
<th>Lateral ventricle volume change (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Mean (s.d.)</td>
<td>Range</td>
<td>Mean (s.d.)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Frazier et al (1996)</td>
<td>21</td>
<td>14.6 (2.1)</td>
<td>6–18</td>
<td>10.2 (1.5)</td>
<td>−9.2</td>
<td>+22.1</td>
</tr>
<tr>
<td>Jacobsen et al (1996)</td>
<td>21</td>
<td>14.6 (2.1)</td>
<td>7–12</td>
<td>10.2 (1.5)</td>
<td>−8.8</td>
<td>−3.3</td>
</tr>
<tr>
<td>Rappoport et al (1997)</td>
<td>16</td>
<td>14.8 (2.4)</td>
<td>9.2–19.1</td>
<td>10.2 (1.7)</td>
<td>−9.8</td>
<td>+28.0</td>
</tr>
<tr>
<td>Friedman et al (1999)</td>
<td>20</td>
<td>14.7 (2.2)</td>
<td>11.3–18.4</td>
<td>12.4 (2.0)</td>
<td>−3.0</td>
<td></td>
</tr>
<tr>
<td>Rappoport et al (1999)</td>
<td>15</td>
<td>13.9 (2.3)</td>
<td>9.2–17.9</td>
<td>10.3 (2.0)</td>
<td>−5.5</td>
<td>−5.5</td>
</tr>
<tr>
<td>Jacobsen et al (1998)</td>
<td>10</td>
<td>15.2 (1.1)</td>
<td>7–12</td>
<td>10.4 (1.7)</td>
<td>−4.06</td>
<td>−1.5</td>
</tr>
<tr>
<td>James et al (1999)</td>
<td>29</td>
<td>16.7 (1.6)</td>
<td>13–20</td>
<td></td>
<td>−1.6</td>
<td>+47.5</td>
</tr>
<tr>
<td>Kumra et al (2000)</td>
<td>44</td>
<td>14.4 (2.3)</td>
<td>6–18</td>
<td></td>
<td>−4.2</td>
<td>−1.5</td>
</tr>
<tr>
<td>Matsumoto et al (2001)</td>
<td>40</td>
<td>15.5 (2.2)</td>
<td>~15</td>
<td></td>
<td>−5.0</td>
<td></td>
</tr>
</tbody>
</table>

1. Follow-up studies.
2. Grey-matter volumes only.

Table 2  Mean verbal, performance and full-scale IQ scores in early-onset cohorts relative to matched normal controls

<table>
<thead>
<tr>
<th>Study</th>
<th>n</th>
<th>Mean age (years)</th>
<th>Age at onset (years)</th>
<th>Verbal Mean (s.d.)</th>
<th>Performance Mean (s.d.)</th>
<th>Full-scale Mean (s.d.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kolvin et al (1971)</td>
<td>33</td>
<td>11.1</td>
<td>5–15</td>
<td>9–11</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Green et al (1984)</td>
<td>23</td>
<td>9.9 (1.6)</td>
<td>6–11</td>
<td>9–10</td>
<td>86 (16.1)</td>
<td>92.9 (16.3)</td>
</tr>
<tr>
<td>Goldberg et al (1988)</td>
<td>30</td>
<td>14.8</td>
<td>13–17</td>
<td>14.5</td>
<td>75.6 (13.9)</td>
<td>71.6 (13.7)</td>
</tr>
<tr>
<td>Assarnow &amp; Ben-Men (1988)</td>
<td>31</td>
<td>10.5 (1.9)</td>
<td>7–13</td>
<td></td>
<td>92.2 (14.3)</td>
<td>87.2 (10.5)</td>
</tr>
<tr>
<td>Russell et al (1989)</td>
<td>35</td>
<td>9.5 (2.0)</td>
<td>4–13</td>
<td>6.9</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Kumra et al (2000)</td>
<td>17</td>
<td>14.4</td>
<td>6–18</td>
<td>10.7</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Matsumoto et al (2001)</td>
<td>40</td>
<td>15.5 (2.2)</td>
<td>~15</td>
<td></td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

Table 3  Demographic characteristics of 33 study participants with early-onset schizophrenia ('cases') and 30 participants in a matched normal control group ('controls')

<table>
<thead>
<tr>
<th></th>
<th>n</th>
<th>Age (years)</th>
<th>Hand preference</th>
<th>Height (cm)</th>
<th>Age at onset (years)</th>
<th>Duration (months)</th>
<th>CPZeq medication (mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Mean (s.d.)</td>
<td>Right/left</td>
<td>Mean (s.d.)</td>
<td>Mean (s.d.)</td>
<td>Mean (s.d.)</td>
<td>Mean (s.d.)</td>
</tr>
<tr>
<td>Cases</td>
<td>33</td>
<td>16.8 (1.4)</td>
<td>25/6</td>
<td>174.3 (10.5)</td>
<td>15.8 (1.4)</td>
<td>12.9 (12.6)</td>
<td>528 (614)</td>
</tr>
<tr>
<td>Male</td>
<td>22</td>
<td>16.8 (1.3)</td>
<td>15/6</td>
<td>179.0 (8.0)</td>
<td>15.9 (1.3)</td>
<td>11.5 (11.4)</td>
<td>438 (345)</td>
</tr>
<tr>
<td>Female</td>
<td>11</td>
<td>16.8 (1.5)</td>
<td>10/0</td>
<td>163.7 (7.6)</td>
<td>15.3 (1.4)</td>
<td>15.6 (14.7)</td>
<td>692 (927)</td>
</tr>
<tr>
<td>Total</td>
<td>33</td>
<td>16.8 (1.4)</td>
<td>25/6</td>
<td>174.3 (10.5)</td>
<td>15.8 (1.4)</td>
<td>12.9 (12.6)</td>
<td>528 (614)</td>
</tr>
<tr>
<td>Controls</td>
<td>30</td>
<td>16.4 (1.7)</td>
<td>27/3</td>
<td>170.4 (8.7)</td>
<td>16.4 (1.7)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>18</td>
<td>16.3 (1.6)</td>
<td>15/3</td>
<td>171.8 (10.0)</td>
<td>16.4 (1.7)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>12</td>
<td>16.4 (1.9)</td>
<td>12/0</td>
<td>168.7 (6.6)</td>
<td>16.4 (1.7)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

CPZeq, chlorpromazine equivalent.
of the temporal lobe was defined as the point where the lateral ventricles divide into frontal and temporal horns. The cerebral hemispheres were separated from the brain-stem at the superior limit of the pons. A more detailed description of these definitions and the methodology is given in Mackay et al (1998). An inter/intrarater reliability study was carried out by three raters. Intraclass correlation coefficients were calculated (Bartko, 1966) and found to be greater than 0.9 for the lateral ventricles, and above 0.8 for temporal lobe and cerebral hemisphere. An index of asymmetry was computed by subtracting the volume of the structure in the left hemisphere (L) from the volume in the right hemisphere (R) and expressing the difference as a percentage of mean volume, i.e. (R − L)/(R + L)/2 × 100. Temporal lobe volume, lateral ventricle volume and asymmetry measures were considered both as absolute values and as proportions of total cerebral hemisphere volume.

### Intentional assessment

Verbal, performance and full-scale IQ data were collected for 28 participants in the ‘cases’ group and 30 in the control group. Five of the original 33 participants did not complete IQ testing, and were not considered in the analysis of IQ. Participants were tested with either the full version of the Wechsler Intelligence Scale for Children – Revised (Wechsler, 1992) or, if they were more than 16 years old, the Wechsler Adult Intelligence Scale – Revised (Wechsler, 1981). Whenever possible, testing was performed in one uninterrupted session. Following testing, all sub-scale scores were transformed into age-scaled scores to render them equivalent. Standard IQ indices were calculated.

**Statistical analysis**

Data were analysed using the Statistical Package for the Social Sciences, version 10 for PC. First, one-way analysis of variance (ANOVA) was performed in order to detect main effects and/or sex interactions in demographic variables. Between-groups comparisons of regional volumes and IQ were performed using the generalised linear model. Multiple analysis of variance (MANOVA) was used to examine structural volume, verbal, performance and full-scale IQ and sub-test differences in IQ performance.

Non-parametric chi-squared analysis was used to determine differences in the distribution of positive and negative asymmetries between groups. All correlations between demographic, treatment and illness-related variables were performed with non-parametric rho connected for multiple comparisons with the Bonferroni test. Given that the normal male brain is significantly larger than the normal female brain (e.g. Gur et al, 1999), and previous studies that show differences in asymmetry between the sexes (e.g. Bear et al, 1986), males and females were examined separately in both volumetric and IQ analyses.

### Results

**Volume measurements**

Absolute mean volumes of the left and right cerebral hemisphere, temporal lobe and ventricle are shown in Table 4. When examined as a group, covarying for sex, participants with schizophrenia had significantly smaller total brain volumes than those in the control group (ANOVA, F=6.37, P=0.01); with an overall difference of 4.5%. In particular, the ‘cases’ group had significantly reduced left (F=7.18, P=0.009) and right (F=5.33, P=0.02) hemisphere volume compared with controls. There was no significant difference in temporal lobe volumes between case and control groups, but a trend to reduced right temporal lobe volume in cases was observed (P=0.08). Volumes of the lateral ventricles were, on average, 20% larger in the cases group, although this difference did not reach statistical significance. No significant difference was found in ventricle volume as a proportion of hemisphere volume between case and control groups.

A significant main effect of sex (F=22.15, P=0.0001) indicated that males as a group had significantly larger brains than females, regardless of diagnosis. Males in the cases group demonstrated a 5.1% reduction in overall brain volume (right plus left hemisphere) relative to males in the control group (MANOVA, F=4.28, P=0.045), whereas females in the case group demonstrated a smaller, 3.5% reduction (P=0.17, NS). MANOVA of individual hemisphere volumes in males and females revealed a significant reduction in left, but not right, hemisphere volume in cases relative to controls in males (F=6.17, P=0.01) but not in females. Males also showed a trend to left temporal lobe reduction compared with male controls (P=0.08).

Overall there was no statistically significant (P<0.05) asymmetry of the cerebral

---

Table 4  Mean volumes (ml) of the left and right hemisphere, lateral ventricle and temporal lobe and mean asymmetry indices for each structure in the group of participants with schizophrenia (‘cases’) and in the control group (‘controls’). Standard deviations are shown in parentheses

<table>
<thead>
<tr>
<th></th>
<th>Hemisphere</th>
<th>Temporal lobe</th>
<th>Lateral ventricle</th>
<th>Asymmetry index</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Left</td>
<td>Right</td>
<td>Left</td>
<td>Right</td>
</tr>
<tr>
<td><strong>Cases</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>547.8 (47.8)</td>
<td>554.0 (52.8)</td>
<td>80.5 (9.8)</td>
<td>80.17 (8.9)</td>
</tr>
<tr>
<td>Female</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>501.0 (33.6)</td>
<td>493.1 (31.8)</td>
<td>72.6 (9.3)</td>
<td>71.1 (8.8)</td>
</tr>
<tr>
<td>Total</td>
<td>532.2 (48.5)</td>
<td>533.7 (54.7)</td>
<td>77.9 (10.2)</td>
<td>77.1 (9.7)</td>
</tr>
<tr>
<td><strong>Controls</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>582.9 (39.6)</td>
<td>579.3 (43.1)</td>
<td>85.5 (7.5)</td>
<td>84.02 (6.1)</td>
</tr>
<tr>
<td>Female</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>522.5 (53.3)</td>
<td>523.8 (52.5)</td>
<td>73.7 (8.6)</td>
<td>74.7 (9.9)</td>
</tr>
<tr>
<td>Total</td>
<td>558.7 (53.8)</td>
<td>557.1 (53.8)</td>
<td>80.8 (9.8)</td>
<td>80.3 (9.8)</td>
</tr>
</tbody>
</table>

H, hemisphere; TL, temporal lobe; LV, lateral ventricle.
hemispheres, temporal lobes or lateral ventricles. However, when sex was entered as
an independent variable, a significant diagnosis × sex interaction was detected in
hemisphere asymmetry after correction for overall brain size (MANOVA, F = 4.39, 
P = .01). Post hoc analyses revealed that the female cases group showed significant
leftward asymmetry of cerebral hemisphere volume (t = 2.28, P = .04), and this was
significantly different from a tendency to rightward asymmetry in the female control
group (F = 4.97, P = .03). No significant asymmetry was detected in males, where
the cases group tended towards rightward asymmetry and the control group towards
leftward asymmetry. In total, 9 out of 11 participants (82%) in the female cases
group demonstrated left greater than right asymmetry, which was present in only 5
out of 12 (41%) of the females in the control group (χ² = 4.5, P < .05). There was
no difference in the proportion of the male cases group relative to the male control
group that showed rightward asymmetry.

IQ analysis
Mean performance IQ, verbal IQ and full-scale IQ scores are shown in Table 5. Both
males and females in the case group showed significant impairments on all three tests
relative to controls (Table 5). Seventy per cent of the early-onset group had IQs below
the mean average range of performance (full-scale IQ less than 90). There was large variability in the average verbal−
performance IQ discrepancy. In the cases group, the average discrepancy was 5.14
IQ points (s.d. = 14.8) compared with −0.16 in controls (s.d. = 17.5) but this was not statistically significant (P = .22).

When the sexes were compared between groups, no significant discrepancy was found between the male case and control
groups (3.21, s.d. = 14.5, and −0.88, s.d. = 19.0, respectively; P = .47). The
female cases group demonstrated a large verbal−performance IQ discrepancy com-
pared with the female control group (9.22, s.d. = 15.4, and 0.83, s.d. = 15.8,
respectively), but this was not statistically significant (P = .23). When individual
Wechsler sub-test scores were examined by MANOVA, significant differences
between case and control groups were found across all 11 sub-tests. The average
sub-test score in the early-onset group was 7.6 (s.d. = 2.0), whereas the control group
average was 3 IQ points higher (10.6, s.d. = 2.4). A 3-point discrepancy in IQ
sub-scale performance is indicative of abnormality (Kaufman, 1990).

Correlation analysis
Age, duration of illness and current medication did not correlate with any of the
structural IQ measures in the cases group. However, earlier age of onset was associated with increasing ventricle volume
as a proportion of total brain volume (r = −0.35, P < .05). There were no statistically significant correlations between IQ
and brain structure measures in the control group as a whole, or for males and females separately. This was also the case for the
combined patient group. However, when the males and females were examined separately, full-scale IQ was significantly correlated with left hemisphere volume (r = 0.47, 
P < .05) in male cases, and verbal IQ was positively correlated with increased
rightward hemispheric asymmetry in female cases (r = .81, P < .01).

DISCUSSION
In this study we examined broad indices of cerebral structure and function in relation
to sex in early-onset schizophrenia. In line with expectations, there was clear evidence
of significant brain volume deficit; a large reduction in intellectual capacity; and a
significant interaction, suggesting that alterations in cerebral asymmetry are sex-
ually dimorphic. Hemisphere volume in males and asymmetry in females were
significantly correlated with IQ in the cases group but not in the control group. The
relationship between brain structure and intellect cannot be understood without
considering the influence of sex.

Volume and IQ
In agreement with previous studies in early-onset schizophrenia (see Tables 1 and 2),
this sample showed an average 4.5% deficit in total brain volume relative to controls,
which was greater on the left than the right, and average full-scale IQ that was close to
the cut-off (80) between low average IQ (IQ = 90–80) and borderline IQ (IQ = 79–66)
ranges of performance. Previous studies have reported greater severity of intellec-
tual disturbance in earlier-onset than in later-onset disease (Yang et al, 1995; Basso
et al, 1997). Our findings corresponded particularly well with those of Matsumoto
et al (2001), who found comparable reductions in brain volume and full-scale IQ in a
cohort of similar age. Together, these findings suggest that alterations in gross
cerebral structure and IQ in early-onset schizophrenia are less severe than those
observed in childhood-onset disease, but greater than those observed in adult-onset
schizophrenia.

Two unexpected findings require further elucidation. First, unlike some pre-
vious studies of normal adults, we did not find significant volume−IQ correlations in
our normal control sample. Most, but not all, studies find modest but significant posi-
tive correlations between overall brain volume (Willerman et al, 1991, 1992;
Andreasen et al, 1993; Wickett et al, 2000) or asymmetry (Yeo et al, 1987; Reiss
et al, 1996) and IQ. It is possible that a larger normal control group would have
revealed comparable volume−IQ correlations. Second, the increase (average 20%)
in ventricular volume in cases relative to controls failed to reach significance, although ventricle-to-brain ratio was inversely related to age at onset. Ventricular enlargement is a robust finding in adult schizophrenia (McCarley et al, 1999) and most studies of early-onset cohorts report increased (James et al, 1999; Kumra et al, 2000; Sowell et al, 2000) and/or progressive enlargement of lateral ventricular volume (Rapport et al, 1997). Our findings indicate considerable variability in the present sample, a finding that is consistent with studies showing variability in ventricular enlargement in first-onset patients (Lieberman et al, 2001; Puri et al, 2001), particularly in the early stages of the illness (Gur et al, 1998; Puri et al, 2001).

**Sex differences**

Our findings provide evidence that changes in brain volume and the relationship to IQ are, to some extent, sexually dimorphic in patients with early-onset schizophrenia. Male, but not female, participants in the cases group had reduced left hemisphere volume and a trend to reduced temporal lobe volume. A significant diagnosis × sex interaction in cerebral hemisphere asymmetry was found such that the tendency for rightward asymmetry in the female control group was reversed to a leftward asymmetry in females with schizophrenia. Males did not show a significant effect, although there was a trend to reversal of the normally observed leftright pattern (i.e. in the opposite direction to females). Sex differences in asymmetry in adults have been previously observed in frontal, temporal and whole brain measurements (Bilder et al, 1994; Highley et al, 1998) and have been shown to interact with age at onset (Highley et al, 1998; Maher et al, 1998; McDonald et al, 2000). Our findings suggest that sex-specific alterations in asymmetry are present in patients with early-onset schizophrenia, consistent with the hypothesis (Crow, 1993, 2000) that a sex-linked determinant of asymmetry has a critical role in the aetiology of schizophrenia.

Correlations between hemisphere volume and IQ measures in this cohort were sex-specific. In the male cases group, reduced left hemisphere volume was correlated with lower full-scale IQ. The reduction in rightward asymmetry in female cases relative to controls was associated with a selective reduction in verbal IQ. In effect, these structure–function relationships were consistent with a body of evidence indicating that sex differences in IQ decline are associated with lateralised cerebral disturbance (see Kaufman, 1990, for review). A simple interpretation of the significant differences and trends in our findings is that in males with schizophrenia the deficits are associated with loss that is relatively selective to the left hemisphere, whereas in females they are associated with a loss that is relatively greater in the right hemisphere. This differs somewhat from the findings of Flbaum et al (1994) who also examined brain structure and IQ in adults with schizophrenia in relation to sex. They reported that the pattern of structure–function correlations in females with schizophrenia was similar to those of female controls, but males with schizophrenia demonstrated no significant structure–function relationships. Further studies are required, but it is noteworthy that Flbaum et al did not examine indices of cerebral asymmetry, where sexual dimorphism in healthy individuals is established (Bear et al, 1986; Barrick et al, 2001).

**Neurodevelopmental antecedents of volume and IQ reduction**

Although the relationship between age, sex and onset of psychosis is complex, our findings are consistent with neurodevelopmental explanations of schizophrenia. Adolescence is a critical stage in cerebral maturation involving substantial volume increases in white matter relative to grey matter (Reiss et al, 1996; Giedd et al, 1997; Courchesne et al, 2000), accompanied by increase in the volume of the lateral ventricles (Giedd et al, 1996, 1997). Furthermore, these changes are influenced by sex, as the extent of grey-matter volume reduction is greater in males than females (Coffey et al, 1998; Raz et al, 1998). Our findings indicate that, in adolescents with schizophrenia, the intellectual deficits depend on an interaction between sex and relative hemispheric development.

**Acknowledgements**

This research project was supported by the Medical Research Council UK (grant G7900348), and SANE. The authors wish to thank Susan James, Gina Clark, Kristin Bohn and Jocasta Webb for their contributions to collecting the sample and assessing and analysing the data.

**References**


- Early-onset schizophrenia is associated with an average 4.5% reduction in brain volume which is greater in males.
- People with early-onset schizophrenia demonstrate significant impairment of intellectual abilities, with the majority of full-scale intelligence quotient scores within the low average to borderline range.
- Significant reduction in intellectual abilities is associated with sexually dimorphic and asymmetrical structural change in early-onset disease consistent with the developmental hypothesis.

Limitations

- The young age range of the participants may limit the general applicability to adult-onset schizophrenia.
- The absence of significant ventricular enlargement is unexpected and may be related to small sample size and high variance in this study.
- Changes in structure are likely to reflect ongoing deviation from normal development. Longitudinal studies including samples in the general population are required.

SIMON L. COLLINSON, DPhil, CLARE E. MACKAY, PhD, POWIC SANE Research Centre, Warneford Hospital, Oxford; ANTHONY C. JAMES, MRCPsych, Highfield Adolescent Unit, Warneford Hospital, Oxford; DIGBY J. QUESTED, MD, Department of Psychiatry, University of Oxford, Warneford Hospital, Oxford; TANIA PHILLIPS, MRCPsych, Highfield Adolescent Unit, Warneford Hospital, Oxford; NEIL ROBERTS, Magnetic Resonance and Image Analysis Research Centre (MARIARC), University of Liverpool, Liverpool; TIMOTHY J. CROW, FMedSci, POWIC SANE Research Centre, University of Oxford, Warneford Hospital, Oxford

Correspondence: Professor T. J. Crow, POWIC SANE Research Centre, University Department of Psychiatry, Warneford Hospital, Oxford OX3 7JX, UK. E-mail: tim.crow@psych.ox.ac.uk

(First received 18 December 2002, final revision 8 April 2003, accepted 22 April 2003)


Brain volume, asymmetry and intellectual impairment in relation to sex in early-onset schizophrenia
SIMON L. COLLINSON, CLARE E. MACKAY, ANTHONY C. JAMES, DIGBY J. QUESTED, TANIA PHILLIPS, NEIL ROBERTS and TIMOTHY J. CROW
BJP 2003, 183:114-120.
Access the most recent version at DOI: 10.1192/bjp.183.2.114

References
This article cites 54 articles, 11 of which you can access for free at:
http://bjp.rcpsych.org/content/183/2/114#BIBL

Reprints/permissions
To obtain reprints or permission to reproduce material from this paper, please write to permissions@rcpsych.ac.uk

You can respond to this article at
/letters/submit/bjprcpsych;183/2/114

Downloaded from
http://bjp.rcpsych.org/ on November 4, 2016
Published by The Royal College of Psychiatrists